#### **Search Criteria**

Report Runtime:	2013-12-19 - 8:07:36 AM			
Range for Initial Receive Date:	1965-01-01 to 2013-08-31			
Product Group	ENERGY DRINKS/BOISSONS ÉNERGISANTES			
Product Role:	Suspect			
Dosage Form:	-All-			
Route of Administration:	-AII-			
Range for Age (years):	-AII-			
Patient Gender:	-AII-			
Case Serious?	-AII-			
Case Outcome:	-AII-			
Report Source:	-All-			
Reporter Type:	-All-			
Domestic or Foreign:	Domestic			

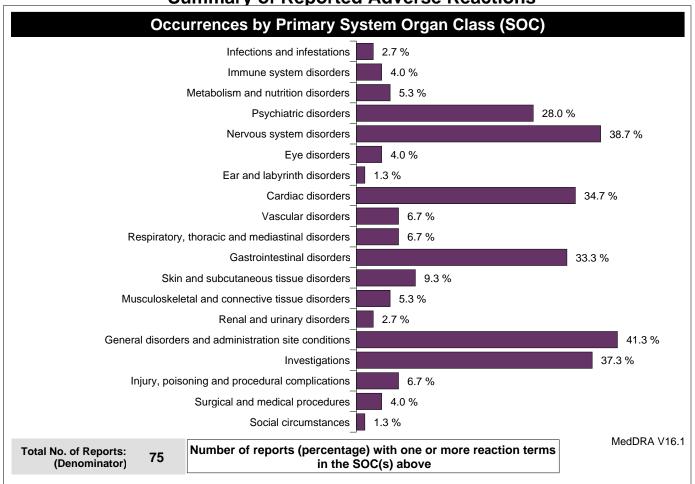
CAVEAT: This summary is based on information from adverse event reports submitted by health professionals and laypersons either directly to Health Canada or via market authorization holders. Each report represents the suspicion, opinion or observation of the individual reporter. The Canada Vigilance Program is a spontaneous reporting system that is suitable to detect signals of potential health product safety issues during the post-market period. The data has been collected primarily by a spontaneous surveillance system in which adverse reactions to health products are reported on a voluntary basis. Under reporting of adverse reactions is seen with both voluntary and mandatory spontaneous surveillance systems. Accumulated case reports should not be used as a basis for determining the incidence of a reaction or estimating risk for a particular product as neither the total number of reactions occurring, nor the number of patients exposed to the health product is known. Because of the multiple factors that influence reporting, quantitative comparisons of health product safety cannot be made from the data. Some of these factors include the length of time a drug is marketed, the market share, size and sophistication of the sales force, publicity about an adverse reaction and regulatory actions. In some cases, the reported clinical data is incomplete and there is not certainty that these health products caused the reported reactions. A given reaction may be due to an underlying disease process or to another coincidental factor. This information is provided with the understanding that the data will be appropriately referenced and used in conjunction with this caveat statement. (10/2007)



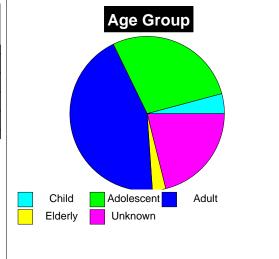
Report Runtime **Product Group:** Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM **NKS/BOISSONS ÉNERGISANTES** 1965-01-01 to 2013-08-31 75 Reports

Canada Vigilance **Summary of Reported Adverse Reactions** 



Age Group							
<b>Child</b> 3 4.00 %							
Adolescent	21	28.00 %					
Adult	33	44.00 %					
Elderly	2	2.67 %					
Unknown	16	21.33 %					



Reason for Seriousness					
Death	3				
LifeThreatening	5				
Hospitalization Required	19				
Disability 4					
Congenital Anomaly	0				
Other Medically Imp Condition	25				

#### **Patient Sex**

Female	17	22.67 %
Male	51	68.00 %
Not specified	6	8.00 %
Unknown	1	1.33 %

The Definition for age groups has been taken from the International Conference on Harmonization (ICH) and have been defined as follows:

- Neonate greater than zero and up to 27 days, inclusively
- Infant greater than 27 days and less than 2 years
  Child greater or equal to 2 and less than 12 years
- Adolescent greater or equal to 12 and less than 19 years
- Adult greater or equal to 19 and less than 65 years
- Elderly greater or equal to 65 years

Serious Reports				
Yes 42				
No	33			



Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

#### **No Duplicate or Linked Reports**

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000128377	0	1999-12-30	1999-12-30	Community		Spontaneous	Other Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:		Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Yes	Other Medically Imp Condition:	

Patient Information

Age	Gender	Height	Weight	Report Outcome
41 Years	Female	168 Centimetres	61 Kilograms	Recovered/resolved

**Product Information** 

Product information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
SOBE - BERRY ENERGY	Suspect	LIQUID	Oral		1 every 1 Day(s)	5 Day(s)
SOBE - CRANBERRY GRAPEFRUIT	Suspect	LIQUID	Oral		every 1 Day(s)	5 Day(s)
SOBE - ORANGE CARROT	Suspect	LIQUID	Oral		every 1 Day(s)	5 Day(s)
SOBE - TROPICAL ENERGY	Suspect	LIQUID	Oral		every 1 Day(s)	5 Day(s)
IMURAN	Concomitant	NOT SPECIFIED	Oral	100 Milligram	every 2 Day(s)	
NEORAL	Concomitant	CAPSULE	Oral	75 Milligram	1 every 1 Day(s)	
PREDNISONE	Concomitant	NOT SPECIFIED	Oral	10 Milligram	every 2 Day(s)	

Redottori iliformation		
MedDRA Preferred Term	MedDRA Version	Duration
Blood creatinine increased	MedDRA V16.1	
Transplant rejection	MedDRA V16.1	



**ETHANOL** 

Santé Canada

# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: VKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000168041	0	2003-10-31	2003-10-31	Other		Spontaneous	Other Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
26 Years	Female			Not recovered/not resolved

Concomitant

NOT SPECIFIED

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BUILL LIQUID VITAMIN	Suspect	LIQUID ORAL				

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Arrhythmia	MedDRA V16.1	
Depressed level of consciousness	MedDRA V16.1	
Dizziness	MedDRA V16.1	
Hypotension	MedDRA V16.1	
Vomiting	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31

75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000172339	0	2004-06-22	2004-06-22	Other		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
Adolescent	Male			Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL LIQUID VITAMIN	Suspect	LIQUID ORAL	Unknown			

Reaction Information

1	MedDRA Preferred Term	MedDRA Version	Duration
	Vomiting	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000175581	0	2004-10-29	2004-10-29	Community		Spontaneous	Other Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disabili	y:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalizatio	ո։	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	

Age	Gender	Height	Weight	Report Outcome
28 Years	Female			Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	3 Dosage forms		
ETHANOL	Concomitant	NOT SPECIFIED	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Dyspnoea	MedDRA V16.1	
Feeling cold	MedDRA V16.1	
Tremor	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000176172	0	2004-11-22	2004-11-22	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number	
No Duplicate or Linked Reports		

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion				
Age	Gender	Heig	ght	Weight	Report Outcome
21 Years	Male				Recovered/resolved

21 Years	Male		Recovered/resolved
Product Informa	ation		

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ETHANOL	Suspect	NOT SPECIFIED	Unknown			
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	500 mL		

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Abdominal pain	MedDRA V16.1	
Blood potassium decreased	MedDRA V16.1	
Vomiting	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000182406	0	2005-08-31	2005-08-31	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:	

Patient Information	tion			
Age	Gender	Height Weight		Report Outcome
35 Years	Male			Not recovered/not resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	250 mL	every 1 Day(s)	1 Month(s)

MedDRA Preferred Term	MedDRA Version	Duration
Insomnia	MedDRA V16.1	
Paranoia	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000190315	0	2005-07-05	2005-07-05	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion				
Age	Gender	Heig	Height Weight		Report Outcome
	Male				Not recovered/not resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect LIQUID ORAL		Oral	750 mL	every 1 Day(s)	

Rodellon information		
MedDRA Preferred Term	MedDRA Version	Duration
Depersonalisation	MedDRA V16.1	
Fatigue	MedDRA V16.1	
Malaise	MedDRA V16.1	
Nausea	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31

75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000190588	0	2005-07-05	2005-07-05	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
	Male			Unknown

**Product Information** 

Product Description	Product Role Dosage Form		Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL		750 mL		

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Depersonalisation	MedDRA V16.1	
Nausea	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000190681	0	2005-09-29	2005-09-29	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number	
No Duplicate or Linked Reports		

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
10 Years	Male			Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MOUNTAIN DEW ENERGY	Suspect	SOLUTION	Unknown	591 mL	1 every 1 Day(s)	

MedDRA Preferred Term	MedDRA Version	Duration
Convulsion	MedDRA V16.1	
Tachycardia	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000194051	0	2005-12-22	2005-12-22	Community		Spontaneous	Pharmacist

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:		Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Yes	Hospitalization:	Other Medically Imp Condition:	

Patient Informat	tion			
Age Gender Hei		Height	Weight	Report Outcome
68 Years	Male			Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MOUNTAIN DEW SUGAR FREE ENERGY DRINK	Suspect	LIQUID	Oral	710 mL		
INSULIN	Concomitant	NOT SPECIFIED				

Reaction information		
MedDRA Preferred Term	MedDRA Version	Duration
Hyperglycaemia	MedDRA V16.1	
Malaise	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: VKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000199896	0	2006-05-30	2006-05-30	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age	ge Gender Hei		nt Weight Report Outcom		
34 Years	Male	183 Centimetres	84 Kilograms	Recovered/resolved	

34 Years	34 Years Male 183 Cer		84 Kilograms	Recovered/resolved
Product Inform	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Oral	473 mL		1 Day(s)

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Hyperhidrosis	MedDRA V16.1	
Nausea	MedDRA V16.1	
Vomiting	MedDRA V16.1	



### **Canada Vigilance Summary of Reported Adverse Reactions**

Report Runtime Initial date of receipt:

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31

**Total Number of Reports:** 75 Reports

Report Information	
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000208519	0	2006-10-06	2006-10-06	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number			
No Duplicate or Linked Reports				

Serious Report?	Death:	Yes	Disability:	Congenital Anomaly:	
Yes	Life Threatening:		Hospitalization:	Other Medically Imp Condition:	

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
15 Years	Male			Death

Product Information	
---------------------	--

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Unknown			
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Arrhythmia	MedDRA V16.1	
Blood pressure increased	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000209393	0	2006-11-16	2006-11-16	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
17 Years	Female	173 Centimetres	64 Kilograms	Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ASSAULT MONSTER ENERGY	Suspect	LIQUID	Oral	473 mL		1 Day(s)
KHAOS MONSTER ENERGY JUICE	Suspect	LIQUID	Unknown	473 mL		1 Day(s)

Reaction Information MedDRA Preferred Term MedDRA Version Duration Chills MedDRA V16.1 Diarrhoea MedDRA V16.1 Dizziness MedDRA V16.1 Hyperhidrosis MedDRA V16.1 Tachycardia MedDRA V16.1 Vomiting MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM
Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000209893	0	2006-11-16	2006-11-16	Other		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Informa	tion			
Age	Gender	Height	Weight	Report Outcome
	Female			Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ETHANOL	Suspect	NOT SPECIFIED	Oral			
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral			-

MedDRA Preferred Term	MedDRA Version	Duration
Diarrhoea	MedDRA V16.1	
Faeces discoloured	MedDRA V16.1	
Insomnia	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000210486	0	2006-12-11	2006-12-11	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Report? Death: Disability:		Congenital Anomaly:			
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes		

Patient Informat	tion				
Age Gender Hei		ght	Weight	Report Outcome	
14 Years	Male				Recovered/resolved

14 Years	Male		Recovered/resolved
Product Informa	ation		

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown	573 mL	every 1 Day(s)	

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Dizziness	MedDRA V16.1	
Feeling abnormal	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	
Hypoaesthesia	MedDRA V16.1	
Palpitations	MedDRA V16.1	



Report Runtime Initial date of receipt:

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31

**Total Number of Reports:** 75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000212817	0	2007-01-23	2007-01-23	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion				
Age Gender Hei		Height	Weight	Report Outcome	
12 Years	Male	150 Centimetres	40 Kilograms	Recovered/resolved	

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
SASKATOON BERRY BEAVER BUZZ JUICE ENERGY	Suspect	LIQUID	Oral	250 mL		
SOBE ARUSH ENERGY DRINK	Suspect	LIQUID	Oral	250 mL		

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Abdominal discomfort	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	
Malaise	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000213842	0	2007-04-10	2007-04-10	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Disability: Congenital Anomaly	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Informat	tion				
Age	Gender	Hei	ght	Weight	Report Outcome
14 Years	Male				Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown			

Reaction information		
MedDRA Preferred Term	MedDRA Version	Duration
Arrhythmia	MedDRA V16.1	
Dry mouth	MedDRA V16.1	
Rash	MedDRA V16.1	
Skin discolouration	MedDRA V16.1	
Speech disorder	MedDRA V16.1	
Tongue oedema	MedDRA V16.1	
Tremor	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000218591	0	2007-05-09	2007-05-09	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number	
No Duplicate or Linked Reports		

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age	Gender	Heig	ght	Weight	Report Outcome
	Male				Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	2 Dosage forms	Once	

Reaction information		
MedDRA Preferred Term	MedDRA Version	Duration
Anxiety	MedDRA V16.1	
Condition aggravated	MedDRA V16.1	
Depression	MedDRA V16.1	
Hyperacusis	MedDRA V16.1	
Insomnia	MedDRA V16.1	
Irritability	MedDRA V16.1	
Restlessness	MedDRA V16.1	



Report Runtime Initial date of receipt:

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31

**Total Number of Reports:** 75 Reports

Report Information
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000218672	0	2007-05-14	2007-05-14	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age Gender Hei		Height	Weight	Report Outcome
				Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RIP IT ENERGY FUEL	Suspect	LIQUID	Oral			

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Chest discomfort	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000220092	0	2007-06-04	2007-06-04	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information

Age	Gender	Height	Weight	Report Outcome
Adolescent				Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Oral	1 Dosage forms		

MedDRA Preferred Term	MedDRA Version	Duration
Arrhythmia	MedDRA V16.1	
Insomnia	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000220931	0	2007-07-05	2007-07-05	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:		
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes	

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
38 Years	Male	163 Centimetres	57 Kilograms	Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RISPERDAL	Suspect	NOT SPECIFIED	Oral	0.5 Milligram	3 every 1 Day(s)	
ROCKSTAR ENERGY DRINK	Suspect	LIQUID				
EFFEXOR NOS	Concomitant	NOT SPECIFIED	Oral			
MULTIVITAMINE(S)	Concomitant	NOT SPECIFIED				
RANITIDINE	Concomitant	NOT SPECIFIED	Oral			
TOPAMAX	Concomitant	NOT SPECIFIED	Oral			
VITAMIN D	Concomitant	NOT SPECIFIED	Oral			
WELLBUTRIN SR	Concomitant	TABLET (EXTENDED-RELEASE)	Oral			

MedDRA Preferred Term	MedDRA Version	Duration
Eye disorder	MedDRA V16.1	
Muscle disorder	MedDRA V16.1	
Muscle twitching	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000221622	0	2007-07-19	2007-07-19	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informa	tion				
Age	Gender	Heigh	t	Weight	Report Outcome
					Recovered/resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
FULL THROTTLE ENERGY DRINK	Suspect	LIQUID	Unknown	1419 mL	every 1 Day(s)	

MedDRA Preferred Term	MedDRA Version	Duration
Disorientation	MedDRA V16.1	
Vomiting	MedDRA V16.1	



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Report Information
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000221635	0	2007-07-24	2007-07-24	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age	Gender	Heig	ght	Weight	Report Outcome
	Male				Not recovered/not resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
GURU ENERGY DRINK - JUICY	Suspect	LIQUID	Oral	473 mL		
TOBACCO	Concomitant	NOT SPECIFIED	Inhalation			

MedDRA Preferred Term	MedDRA Version	Duration
Fatigue	MedDRA V16.1	
Headache	MedDRA V16.1	
Malaise	MedDRA V16.1	
Palpitations	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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eports: 75 Reports

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000222275	0	2007-07-11	2007-07-11	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Inform	nation			
Age	Gender	Heigh	t Weight	Report Outcome
	Female			Recovered/resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Headache	MedDRA V16.1	
Nausea	MedDRA V16.1	



### Canada Vigilance Summary of Reported Adverse Reactions

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000225597	0	2007-12-03	2007-12-03	Community		Spontaneous	Other Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age	Gender	Height		Weight	Report Outcome
	Male				Recovered/resolved

	Product Information						
[	Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration

ETHANOL	Suspect	NOT SPECIFIED			
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown		
`					

#### Reaction Information MedDRA Preferred Term MedDRA Version Duration Amnesia MedDRA V16.1 Discomfort MedDRA V16.1 Malaise MedDRA V16.1 Mental disorder MedDRA V16.1 Speech disorder MedDRA V16.1 Vomiting MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000225870	0	2007-11-27	2007-11-27	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion				
Age	Gender	Height		Weight	Report Outcome
	Female				Recovered/resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
GREENS + EXTRA ENERGY	Suspect	POWDER	Oral	13 Gram	1 every 1 Day(s)	4 Day(s)
SYNTHROID	Concomitant	NOT SPECIFIED	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Dysgeusia	MedDRA V16.1	
Fatigue	MedDRA V16.1	
Hypoaesthesia	MedDRA V16.1	
Nausea	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000226716	0	2007-12-21	2007-12-21	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
30 Years		178 Centimetres	70 Kilograms	Recovered/resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Oral		1 every 1 Day(s)	

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Palpitations	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000307989	0	2008-07-04	2008-07-04	Community		Spontaneous	Pharmacist

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information

Age	Gender	Height	Weight	Report Outcome
25 Years	Male	173 Centimetres	160 Pounds	Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	1 Dosage forms		

MedDRA Preferred Term	MedDRA Version	Duration
Atrial septal defect repair	MedDRA V16.1	
Monoplegia	MedDRA V16.1	
Speech disorder	MedDRA V16.1	
Transient ischaemic attack	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000308238	0	2008-07-08	2008-07-08	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

P	atient Informa	tion				
	Age	Gender	Hei	ght	Weight	Report Outcome
	Adolescent	Not specified				Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown			

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Palpitations	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000310349	3	2008-08-20	2010-03-23	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Yes	Disability:	Congenital Anomaly:	
Yes	Life Threatening:		Hospitalization:	Other Medically Imp Condition:	

Patient Information

Age	Gender	Height	Weight	Report Outcome
15 Years	Male	71 Inches	170 Pounds	Death

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID	Oral	250 mL	Once	Once

MedDRA Preferred Term	MedDRA Version	Duration
Arrhythmia	MedDRA V16.1	
Blood caffeine	MedDRA V16.1	
Circulatory collapse	MedDRA V16.1	
Sudden cardiac death	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000310907	0	2008-09-08	2008-09-08	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
17 Years	Male			Unknown

17 Years	Male		Unknown
Product Informs	ation		

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	3 Dosage forms	1 every 1 Day(s)	

Reaction Information			
	MedDRA Preferred Term	MedDRA Version	Duration
Insomnia		MedDRA V16.1	



ROCKSTAR ENERGY DRINK

### **Canada Vigilance Summary of Reported Adverse Reactions**

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**Therapy Duration** 

1 Day(s)

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000311394	0	2008-09-18	2008-09-18	Community		Spontaneous	Pharmacist

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:		Disability:		Congenital Anomaly:	
Yes	Life Threatening:	Yes	Hospitalization:	Yes	Other Medically Imp Condition:	

Patient Information	tion			
Age	Age Gender Hei		Weight	Report Outcome
20 Years	Female	150 Centimetres	50 Kilograms	Unknown

Suspect

LIQUID

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	

**Reaction Information MedDRA Preferred Term MedDRA Version** Duration MedDRA V16.1 Blood creatine phosphokinase increased Endotracheal intubation MedDRA V16.1 Injection site erythema MedDRA V16.1 Injection site pruritus MedDRA V16.1 Laryngeal oedema MedDRA V16.1 Pharyngeal oedema MedDRA V16.1 MedDRA V16.1 Rash Respiratory distress MedDRA V16.1 Skin test positive MedDRA V16.1 Stridor MedDRA V16.1 Swollen tongue MedDRA V16.1 Throat irritation MedDRA V16.1 Urticaria MedDRA V16.1

Unknown



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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000311650	0	2008-09-19	2008-09-19	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability: Ye	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:

Patient Information	tion				
Age	Gender	Hei	ght	Weight	Report Outcome
46 Years	Male				Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Oral			
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Oral	1 Dosage forms	8 every 1 Day(s)	3 Month(s)

# Reaction Information MedDRA Preferred Term MedDRA Version Duration Dependence MedDRA V16.1 MedDRA V16.1 Drug withdrawal syndrome MedDRA V16.1 MedDRA V16.1 Fatigue MedDRA V16.1 MedDRA V16.1 Weight decreased MedDRA V16.1 MedDRA V16.1



### Canada Vigilance Summary of Reported Adverse Reactions

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000312304	1	2008-09-11	2010-06-07	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Informat	tion			
Age	Age Gender Hei		Weight	Report Outcome
16 Years	Male	152 Centimetres	100 Pounds	Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	650 mL	Once	Once
STRATTERA	Concomitant	CAPSULF	Unknown			

# MedDRA Preferred Term MedDRA Version Duration Electrocardiogram abnormal MedDRA V16.1 Heart rate increased MedDRA V16.1 Malaise MedDRA V16.1 Palpitations MedDRA V16.1



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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000312726	0	2008-10-20	2008-10-20	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number			
No Duplicate or Linked Reports				

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informa	tion			
Age	Gender	Height	Weight	Report Outcome
13 Years	Female			Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Unknown	473 mL	5 every 2 Day(s)	2 Day(s)

MedDRA Preferred Term	MedDRA Version	Duration
Agitation	MedDRA V16.1	
Hyperkinesia	MedDRA V16.1	
Impulsive behaviour	MedDRA V16.1	



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Report Information	
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000313337	0	2008-11-04	2008-11-04	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number				
No Duplicate or Linked Reports					

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age Gender Hei		ght	Weight	Report Outcome	
	Male				Unknown

Product Information

Product Description	Product Role Dosage Form		Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	1 Dosage forms	2 every 1 Day(s)	

reaction information		
MedDRA Preferred Term	MedDRA Version	Duration
Chest pain	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	
Palpitations	MedDRA V16.1	
Syncope	MedDRA V16.1	



### Canada Vigilance Summary of Reported Adverse Reactions

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000315341	0	2008-12-15	2008-12-15	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
24 Years	Female			Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral			
ETHANOL	Concomitant	NOT SPECIFIED	Oral			

#### Reaction Information MedDRA Preferred Term MedDRA Version Duration Anxiety MedDRA V16.1 Fatigue MedDRA V16.1 Insomnia MedDRA V16.1 Irritability MedDRA V16.1 Palpitations MedDRA V16.1 Psychomotor hyperactivity MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000315734	0	2008-12-29	2008-12-29	Community		Spontaneous	Consumer Or Other Non Health Professional

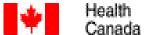
Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:		Disability:		Congenital Anomaly:
Yes	Life Threatening:	Yes	Hospitalization:	Yes	Other Medically Imp Condition:

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
22 Years	Female	165 Centimetres	126 Pounds	Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Oral	660 mL	1 every 1 Day(s)	1 Day(s)
CONTRACEPTIVES	Concomitant	NOT SPECIFIED	Unknown			
REACTINE	Concomitant	NOT SPECIFIED	Unknown			

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Cold sweat	MedDRA V16.1	
Diarrhoea	MedDRA V16.1	
Disturbance in attention	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	
Palpitations	MedDRA V16.1	
Tremor	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000316215	1	2009-01-07	2009-08-21	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:		Disability: Yes		Congenital Anomaly:		
Yes	Life Threatening:	Yes	Hospitalization:	Yes	Other Medically Imp Condition:	Yes	

Patient Information	tion					
Age Gender Hei		Height	Weight	Report Outcome		
18 Years	Male	67 Inches	125 Pounds	Not recovered/not resolved		

Product Information							
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration	
RED BULL ENERGY DRINK	Suspect	LIQUID	Unknown	250 mL	2 every 1 Day(s)		
ETHANOL	Concomitant	NOT SPECIFIED	Unknown				
MARIJUANA	Concomitant	NOT SPECIFIED	Unknown				

Reaction Information **MedDRA Preferred Term MedDRA Version** Duration Amnesia MedDRA V16.1 Blindness unilateral MedDRA V16.1 Cardio-respiratory arrest MedDRA V16.1 Eve movement disorder MedDRA V16.1 Hemiplegia MedDRA V16.1 Hypoxic-ischaemic encephalopathy MedDRA V16.1 Malaise MedDRA V16.1 MedDRA V16.1 Muscle spasms Muscular weakness MedDRA V16.1 MedDRA V16.1 Myopathy toxic **Palpitations** MedDRA V16.1 Pneumonia MedDRA V16.1 MedDRA V16.1 Speech disorder Torsade de pointes MedDRA V16.1 Troponin increased MedDRA V16.1 Ventricular fibrillation MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000316823	0	2009-01-21	2009-01-21	Hospital		Spontaneous	Pharmacist

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability: Ye	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:

Patient Informat	tion				
Age	Gender	Heig	ht	Weight	Report Outcome
68 Years	Male				Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ELECTROLYTE STAMINA TABLETS	Suspect	TABLET	Unknown	10 Dosage forms	1 every 1 Day(s)	
GREENS + EXTRA ENERGY	Suspect	POWDER	Unknown	1 Dosage forms	1 every 1 Day(s)	
RECOVERY	Suspect	POWDER	Unknown	3 Teaspoonful	1 every 1 Day(s)	
VIGRX PLUS	Suspect	TABLET	Unknown	2 Dosage forms	1 every 1 Day(s)	

Troublion information			
·	ledDRA Preferred Term	MedDRA Version	Duration
Biopsy kidney abnormal		MedDRA V16.1	
Nephropathy toxic		MedDRA V16.1	
Renal failure		MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000316861	0	2009-01-16	2009-01-16	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
25 Years	Male	70 Inches	190 Pounds	Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
JOLT ENERGY	Suspect	LIQUID ORAL	Unknown			
MONSTER ENERGY	Suspect	LIQUID	Unknown			
RAGE LIQUID ENERGY	Suspect	LIQUID ORAL	Unknown			
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Disturbance in attention	MedDRA V16.1	
Headache	MedDRA V16.1	
Hypersomnia	MedDRA V16.1	
Irritability	MedDRA V16.1	
Reflux gastritis	MedDRA V16.1	
Withdrawal syndrome	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000319579	0	2009-03-09	2009-03-09	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age Gender Heig		ght	ht Weight Report Outcome		
	Male		•		Not recovered/not resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	1 Dosage forms	1 every 1 Day(s)	2 Year(s)
MONSTER ENERGY	Suspect	LIQUID	Oral	1 Dosage forms	2 every 1 Day(s)	2 Year(s)

Nousier in emailer		
MedDRA Preferred Term	MedDRA Version	Duration
Dependence	MedDRA V16.1	
Faecal incontinence	MedDRA V16.1	
Heart rate irregular	MedDRA V16.1	
Metabolic disorder	MedDRA V16.1	
Renal pain	MedDRA V16.1	
Urine abnormality	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

75 Reports

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000321253	1	2009-03-17	2009-04-06	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion					
Age Gender Hei		Height	Weight	Report Outcome		
22 Years	Male	68 Inches	190 Pounds	Unknown		

22 Years	22 Years Male 68 In		190 Pounds	Unknown	
Product Informa	ation				

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Unknown	500 mL	1 every 3 Day(s)	366

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Disturbance in attention	MedDRA V16.1	
Dizziness	MedDRA V16.1	
Dyspnoea	MedDRA V16.1	
Heart rate irregular	MedDRA V16.1	
Hyperhidrosis	MedDRA V16.1	
Nausea	MedDRA V16.1	
Pyrexia	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000322030	0	2009-04-20	2009-04-20	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

<b>Patient Informa</b>	tion			
Age	Gender	Height	Weight	Report Outcome
38 Years	Male	180 Centimetres	86 Kilograms	Unknown

Product Information	

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown			

#### Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Heart rate increased	MedDRA V16.1	
Heart rate irregular	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000322689	0	2009-04-30	2009-04-30	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

#### Patient Information

Age	Gender	Height	Weight	Report Outcome
31 Years	Male	177 Centimetres	150 Pounds	Unknown

#### **Product Information**

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	355 mL	Once	Once

MedDRA Preferred Term	MedDRA Version	Duration
Heart rate increased	MedDRA V16.1	
Palpitations	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

eports: 75 Reports

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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000322939	0	2009-05-04	2009-05-04	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
30 Years	Female			Unknown

Product Information

	Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
- {	RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown			

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Heart rate increased	MedDRA V16.1	
Heart rate irregular	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000323383	0	2009-04-29	2009-04-29	Community		Spontaneous	Pharmacist

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
23 Years	Male	70 Inches	190 Pounds	Recovered/resolved

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
TURBO ENERGY DRINK	Suspect	LIQUID	Unknown		1 every 1 Week(s)	

MedDRA Preferred Term	MedDRA Version	Duration
Back pain	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

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Product Group: NKS/BOISSONS ÉNERGISANTES
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000323401	0	2009-05-05	2009-05-05	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition: Ye	es

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
17 Years	Female	67 Inches	130 Pounds	Not recovered/not resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Unknown			6 Month(s)
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	2 Dosage forms		6 Month(s)

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Convulsion	MedDRA V16.1	
Disturbance in attention	MedDRA V16.1	
Educational problem	MedDRA V16.1	
Fatigue	MedDRA V16.1	
Memory impairment	MedDRA V16.1	
Migraine	MedDRA V16.1	
Palpitations	MedDRA V16.1	



**Reaction Information** 

### **Canada Vigilance Summary of Reported Adverse Reactions**

Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000324299	0	2009-05-25	2009-05-25	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informa	tion			
Age	Gender	Height	Weight	Report Outcome
	Unknown			Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
JAVA MONSTER ENERGY DRINK	Suspect	LIQUID	Unknown			

MedDRA Preferred Term **MedDRA Version** Duration Stomatitis MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

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date of receipt: 1965-01-01 to 2013-08-31

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000324449	0	2009-06-01	2009-06-01	Hospital		Spontaneous	Pharmacist

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:

Patient Information				
Age	Gender	Height	Weight	Report Outcome
29 Years	Male	74 Inches	243 Pounds	Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ETHANOL	Suspect	NOT SPECIFIED	Unknown			
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	4 Dosage forms	Total	
IBUPROFEN	Concomitant	NOT SPECIFIED	Oral			

# Reaction Information MedDRA Version Duration Atrial fibrillation MedDRA V16.1 Blood alcohol increased MedDRA V16.1 Heart rate increased MedDRA V16.1 Hypertension MedDRA V16.1 Palpitations MedDRA V16.1 MedDRA V16.1



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000325467	0	2009-06-19	2009-06-19	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Informat	tion			
Age Gender Hei		Height	Weight	Report Outcome
15 Years	Male			Not recovered/not resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Oral			

TOWARD THE CONTRACTOR		
MedDRA Preferred Term	MedDRA Version	Duration
Asthenia	MedDRA V16.1	
Loss of consciousness	MedDRA V16.1	
Pallor	MedDRA V16.1	
Palpitations	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000326483	2	2009-07-14	2010-07-21	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	-	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Yes	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
18 Years	Male	73 Inches	145 Pounds	Recovered/resolved

**Product Information** 

**Reaction Information** 

Convulsion

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	375 mL	Once	

**MedDRA Preferred Term** MedDRA Version Duration MedDRA V16.1 Accidental overdose Amnesia MedDRA V16.1 Blood magnesium increased MedDRA V16.1 Blood phosphorus decreased MedDRA V16.1

MedDRA V16.1

MedDRA V16.1 Dehydration Disorientation MedDRA V16.1 Dizziness MedDRA V16.1 Fall MedDRA V16.1 Feeling abnormal MedDRA V16.1

Headache MedDRA V16.1 Loss of consciousness MedDRA V16.1 Neutrophil count increased MedDRA V16.1 MedDRA V16.1 Scratch MedDRA V16.1 Tongue injury White blood cell count increased MedDRA V16.1



Report Runtime Initial date of receipt: **Total Number of Reports:** 

2013-12-19 - 8:07:36 AM Product Group: NKS/BOISSONS ÉNERGISANTES 1965-01-01 to 2013-08-31 75 Reports

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000329550	0	2008-03-07	2008-03-07	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:	

Patient Informa	tion			
Age	Gender	Height	Weight	Report Outcome
12 Years	Female		80 Pounds	Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	8 Dosage forms	1 every 1 Day(s)	

MedDRA Preferred Term	MedDRA Version	Duration
Gastric lavage	MedDRA V16.1	
Intentional overdose	MedDRA V16.1	



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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000331450	0	2009-10-15	2009-10-15	Hospital		Spontaneous	Pharmacist

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:

Patient Information	tion				
Age Gender Hei		Height	nt Weight Report Outcome		
21 Years	Female	173 Centimetres	94 Kilograms	Recovered/resolved	

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ETHANOL	Suspect	NOT SPECIFIED	Unknown	6 Dosage forms		1 Day(s)
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown			1 Day(s)
CHLORPROMAZINE	Concomitant	NOT SPECIFIED	Unknown			
CIPRALEX	Concomitant	TABLET	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Atrial fibrillation	MedDRA V16.1	
Blood bicarbonate decreased	MedDRA V16.1	
Blood pH increased	MedDRA V16.1	
Palpitations	MedDRA V16.1	
PCO2 decreased	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

Report Runtime Product Group: Initial date of receipt: Total Number of Reports:

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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000332689	0	2009-11-09	2009-11-09	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion				
Age Gender Hei		Height	nt Weight Report Outcome		
30 Years	Female	167 Centimetres	56 Kilograms	Recovered/resolved	

30 Years	Female	167 Centimetres	56 Kilograms	Recovered/resolved
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	1 Dosage forms	Once	Once
· · · · · · · · · · · · · · · · · · ·						

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Feeling abnormal	MedDRA V16.1	
Flushing	MedDRA V16.1	
Palpitations	MedDRA V16.1	



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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000333118	0	2009-11-17	2009-11-17	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number	
No Duplicate or Linked Reports		

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion			
Age Gender Hei		Height	Weight	Report Outcome
19 Years	Male	71 Inches	165 Pounds	Unknown

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	473 mL	Once	Once
DEXEDRINE	Concomitant	NOT SPECIFIED	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Eyelid function disorder	MedDRA V16.1	
Hypersomnia	MedDRA V16.1	
Initial insomnia	MedDRA V16.1	



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000337976	1	2010-02-15	2010-07-21	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age Gender Hei		Height	Weight	Report Outcome
32 Years	Male	170 Centimetres	184 Pounds	Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL SUGAR FREE	Suspect	LIQUID	Oral	200 mL	Once	1 Day(s)
ACIDOPHYLUS	Concomitant	NOT SPECIFIED	Unknown			
DIOVAN	Concomitant	NOT SPECIFIED	Unknown			
LIPITOR	Concomitant	TABLET	Unknown			
NOVORAPID	Concomitant	NOT SPECIFIED	Unknown	·		

**Reaction Information** MedDRA Preferred Term **MedDRA Version** Duration Blood glucose fluctuation MedDRA V16.1 Blood glucose increased MedDRA V16.1 Chest pain MedDRA V16.1 Heart rate increased MedDRA V16.1



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000341621	1	2010-04-15	2010-05-11	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
19 Years	Male	73 Inches	190 Pounds	Recovered/resolved

19 Years	Male	73 Inches	190 Pounds	Recovered/resolved
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR RECOVERY	Suspect	LIQUID ORAL	Oral	473 mL	Once	
Reaction Information						

Reaction information		
MedDRA Preferred Term	MedDRA Version	Duration
Erythema	MedDRA V16.1	1 Day(s)
Fatigue	MedDRA V16.1	1 Day(s)
Nausea	MedDRA V16.1	1 Day(s)
Pruritus	MedDRA V16.1	1 Day(s)
Pyrexia	MedDRA V16.1	1 Day(s)
Swelling face	MedDRA V16.1	1 Day(s)
Urticaria	MedDRA V16.1	1 Day(s)



Report Runtime Initial date of receipt: **Total Number of Reports:** 

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000343925	0	2010-05-21	2010-05-21	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

<b>Patient Informa</b>	ation			
Age	Gender	Height	Weight	Report Outcome
	Not specified			Recovered/resolved

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	2 Dosage forms		

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Emotional distress	MedDRA V16.1	

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000347446	0	2010-07-16	2010-07-16	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Informat	tion			
Aae	Gender	Heiaht	Weiaht	Report Outcome
57 Years	Male	170 Centimetres	71 Kilograms	Recovered/resolved

Aae	Gender	Heiaht	Weiaht	Report Outcome
57 Years	Male	170 Centimetres	71 Kilograms	Recovered/resolved
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Oral	650 mL	Once	Once

Reaction	Informati	ion
Neaction	minomia	1011

MedDRA Preferred Term	MedDRA Version	Duration
Feeling jittery	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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Report Runtime 2013-12-19 - 8:07:36 AM

Product Group: NKS/BOISSONS ÉNERGISANTES
date of receipt: 1965-01-01 to 2013-08-31

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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000351921	0	2010-09-23	2010-09-23	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:		Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Yes	Other Medically Imp Condition:	Yes

<b>Patient Informat</b>	tion			
Age	Gender	Height	Weight	Report Outcome
21 Years	Female			Unknown

Product Information

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK	Suspect	LIQUID	Unknown			

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Hypersensitivity	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000351922	0	2010-09-23	2010-09-23	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information

Age	Gender	Height	Weight	Report Outcome
10 Years	Male			Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
ROCKSTAR ENERGY DRINK 710ML	Suspect	LIQUID ORAL	Oral	1 Litres	Total	

Ł	MedDRA Preferred Term	MedDRA Version	Duration
[	Vomiting	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000351923	0	2010-09-23	2010-09-23	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	-	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Yes	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
35 Years	Male			Unknown

Product Information

	Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
R	OCKSTAR RECOVERY	Suspect	LIQUID ORAL	Unknown			

Reaction Information

MedDRA Preferred Term	MedDRA Version	Duration
Hypersensitivity	MedDRA V16.1	

#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000352663	0	2010-10-04	2010-10-04	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information

Age	Gender	Height	Weight	Report Outcome
17 Years	Male	187 Centimetres	95 Kilograms	Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	374 mL	Once	Once

MedDRA Preferred Term	MedDRA Version	Duration
Blood pressure increased	MedDRA V16.1	
Headache	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	
Restlessness	MedDRA V16.1	



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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000353286	0	2010-10-14	2010-10-14	MAH		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age	Gender	Hei	ght	Weight	Report Outcome
	Male				Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
FULL THROTTLE	Suspect	LIQUID	Unknown	1 Dosage forms		

MedDRA Preferred Term	MedDRA Version	Duration
Diarrhoea	MedDRA V16.1	
Vomiting	MedDRA V16.1	



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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000353507	1	2010-10-17	2010-12-10	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
57 Years	Male	178 Centimetro	es 205 Pounds	Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL 355 ML ENERGY DRINK	Suspect	LIQUID	Oral	1 Dosage forms	Once	Once

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Amnesia	MedDRA V16.1	
Cerebral ischaemia	MedDRA V16.1	
Palpitations	MedDRA V16.1	
Transient global amnesia	MedDRA V16.1	



# Canada Vigilance Summary of Reported Adverse Reactions

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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000360030	1	2011-01-20	2011-07-05	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Informa	tion			
Age Gender Hei		Height	Weight	Report Outcome
35 Years	Male	182 Centimetres	180 Pounds	Recovered/resolved

35 Years	Male	182 Centimetres	180 Pounds	Recovered/resolved
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	473 mL	Once	Once

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Chest discomfort	MedDRA V16.1	1 Month(s)
Dyspnoea	MedDRA V16.1	1 Month(s)
Heart rate irregular	MedDRA V16.1	1 Month(s)
Palpitations	MedDRA V16.1	1 Month(s)



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Report Information	
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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000360557	1	2011-01-26	2011-02-03	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number			
No Duplicate or Linked Reports				

Serious Report?	Death:	N/A	Disability:	N/A	Congenital Anomaly:	N/A
No	Life Threatening:	N/A	Hospitalization:	N/A	Other Medically Imp Condition:	N/A

Patient Information	tion				
Age Gender Hei		ght	Weight	Report Outcome	
	Male				Recovered/resolved

Age	Gender	Height	Weight	Report Outcome
	Male			Recovered/resolved
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
FULL THROTTLE	Suspect	LIQUID	Oral	1 Dosage forms		

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Dyskinesia	MedDRA V16.1	
Feeling abnormal	MedDRA V16.1	
Heart rate increased	MedDRA V16.1	



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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000362024	1	2011-02-15	2011-03-01	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
42 Years	Male	74 Inches	220 Pounds	Recovered/resolved with sequelae

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	473 mL	5 every 1 Week(s)	751
MONSTER ENERGY	Suspect	LIQUID	Oral	946 mL	Once	Once
COFFEE	Concomitant		Unknown			

#### Reaction Information MedDRA Preferred Term MedDRA Version Duration Convulsion MedDRA V16.1 1 Day(s) Grand mal convulsion MedDRA V16.1 1 Day(s) Spinal compression fracture MedDRA V16.1



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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000368520	1	2011-05-10	2011-07-13	Hospital		Spontaneous	Physician

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition:	

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
15 Years	Male		110 Pounds	Recovered/resolved

15 Years	Male	110 Pounds	Recovered/resolved
Product Informa	ation		

Product Description	Product Role	Dosage Form		Route	Dosing	Frequency	Therapy Duration
MONSTER NITROUS ANTI-GRAVITY ENERGY	Suspect	LIQUID	Oral		1 Dosage forms	Once	Once
Reaction Information							
MedDRA Preferred Term				MedDRA Ve	ersion		Duration
Blood creatine phosphokinase increased			MedDRA \	/16.1			

MedDRA Preferred Term	MedDRA Version	Duration
Blood creatine phosphokinase increased	MedDRA V16.1	
Chest pain	MedDRA V16.1	1 Day(s)
Electrocardiogram ST segment elevation	MedDRA V16.1	
Feeling abnormal	MedDRA V16.1	
Malaise	MedDRA V16.1	
Pericarditis	MedDRA V16.1	
Syncope	MedDRA V16.1	
Troponin I increased	MedDRA V16.1	10 Day(s)
Viral upper respiratory tract infection	MedDRA V16.1	
Vomiting	MedDRA V16.1	



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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000372282	2	2011-06-24	2011-09-07	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
8 Years	Male	54 Inches	80 Pounds	Recovered/resolved

**Product Information** 

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY ABSOLUTELY ZERO	Suspect	LIQUID	Unknown		Once	Once
CLARITIN KIDS	Concomitant	SYRUP	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Blood caffeine increased	MedDRA V16.1	
Dyspnoea	MedDRA V16.1	
Palpitations	MedDRA V16.1	
Sleep disorder	MedDRA V16.1	



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Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000378905	0	2011-09-15	2011-09-15	Community		Spontaneous	Consumer Or Other Non Health Professional

Record Type	Link Aer Number
No Duplicate or Linked Reports	

Serious Report?	Death:	Yes	Disability:	-	Congenital Anomaly:
Yes	Life Threatening:		Hospitalization:		Other Medically Imp Condition:

Patient Information	tion			
Age	Gender	Height	Weight	Report Outcome
18 Years	Male			Death

Age	Gender	Height	Weight	Report Outcome
18 Years	Male			Death
Product Informa	ation			

Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
RED BULL ENERGY DRINK	Suspect	LIQUID	Oral	250 mL	4 every 1 Hour(s)	1 Hour(s)

Reaction Information		
MedDRA Preferred Term	MedDRA Version	Duration
Delirium	MedDRA V16.1	
Hallucination	MedDRA V16.1	
Hypoglycaemia	MedDRA V16.1	
Poisoning	MedDRA V16.1	



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#### Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000387213	0	2011-11-19	2011-11-19	Community		Spontaneous	Other Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:		Disability:	Yes	Congenital Anomaly:	
Yes	Life Threatening:	Yes	Hospitalization:	Yes	Other Medically Imp Condition:	Yes

Patient Informat	tion			
Age	Gender	Height	Weight	Report Outcome
18 Years	Male	185 Centimetres	81 Kilograms	Unknown

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
5-HOUR ENERGY	Suspect	LIQUID	Unknown			
NOS HIGH PERFORMANCE ENERGY DRINK	Suspect	LIQUID ORAL	Unknown	3 Dosage forms		
RED BULL ENERGY DRINK	Suspect	LIQUID ORAL	Oral	3 Dosage forms		1 Week(s)
RED RAIN	Suspect	LIQUID	Unknown			

MedDRA Preferred Term	MedDRA Version	Duration
Concussion	MedDRA V16.1	
Delusion	MedDRA V16.1	
Disturbance in attention	MedDRA V16.1	
Hallucination	MedDRA V16.1	
Multiple fractures	MedDRA V16.1	



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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000480736	0	2012-11-28	2012-11-28	Hospital		Spontaneous	Other Health Professional

Record Type	Link Aer Number		
No Duplicate or Linked Reports			

Serious Report?	Death:	Disability:	Congenital Anomaly:	
Yes	Life Threatening:	Hospitalization:	Other Medically Imp Condition:	Yes

Patient Information	tion				
Age	Gender	Hei	ght	Weight	Report Outcome
24 Years	Male			174 Pounds	Recovered/resolved

ſ		. D	Dundred Dala	D		D 4 -
	Product Informa	ation				
١	24 Years	Male	174 Pour	nds Recovere	ed/resolved	
1	24 Years	Male	174 Pour	nds Recovere	ed/resolved	

200 1	
nknown 300 mL	
nknown	

MedDRA Preferred Term		MedDRA Version	Duration
Nausea		MedDRA V16.1	
Tremor		MedDRA V16.1	
Vomiting		MedDRA V16.1	



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Report Information

Aer No	Version No.	Initial Rec. Date	Latest Rec. Date	Report Source	MAH Number	Type of Report	Reporter Type
000520801	0	2013-05-01	2013-05-01	Hospital		Spontaneous	Pharmacist

Record Type	Link Aer Number	
No Duplicate or Linked Reports		

Serious Report?	Death:	Disability:	Congenital Anomaly:
Yes	Life Threatening:	Hospitalization: Yes	Other Medically Imp Condition: Yes

Patient Informa	tion			
Age	Gender	Height	Weight	Report Outcome
19 Years	Male	61 Inches	173 Pounds	Recovered/resolved

Product Information						
Product Description	Product Role	Dosage Form	Route	Dosing	Frequency	Therapy Duration
MONSTER ENERGY	Suspect	LIQUID	Oral	1 Dosage forms	1 every 1 Day(s)	1 Week(s)
SIX STAR MUSCLE PROFESSIONAL STRENGTH MUSCLE BUILDING MILKSHAKE	Suspect	POWDER	Oral	2 Dosage forms	2 every 1 Day(s)	1 Week(s)
SIX STAR PROFESSIONAL STRENGTH CREATINE X3 POWDER	Suspect	POWDER	Oral	2 Dosage forms	1 every 1 Day(s)	1 Week(s)

MedDRA Preferred Term	MedDRA Version	Duration
Drug interaction	MedDRA V16.1	
Rhabdomyolysis	MedDRA V16.1	