

Ontario Drug Benefit Formulary/Comparative Drug Index

Edition 43

Summary of Changes – December 2024 Effective December 30, 2024

Drug Programs Policy and Strategy Branch Health Programs and Delivery Division Ministry of Health

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New Single Source Products

Generic Name: AFLIBERCEPT

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02545004	Eylea HD	8mg/0.07mL	Inj Sol-0.07mL Vial Pk	BAH	1250.0000/Vial

Reason For Use Code and Clinical Criteria

Code 694

For the treatment of patients with neovascular (wet) age-related macular degeneration (AMD) in a treatment-naive eye.

Initial diagnosis should be confirmed by an appropriate diagnostic procedure and administration should be done by a qualified ophthalmologist experienced in intravitreal injections.

Patients receiving concurrent administration of other anti-VEGF intravitreal injections are not eligible for reimbursement.

Treatment with anti-VEGF agents should only be continued in patients who maintain adequate response to therapy.

Coverage will be provided for patients responding to therapy with another anti-VEGF agent who switch to Eylea HD. Coverage will not be provided for patients who have failed to respond to other anti-VEGF agents.

Recommended Dose: Treatment should be initiated with a monthly intravitreal injection for the first 3 consecutive doses, followed by one injection every 8 to 16 weeks.

LU Authorization Period: 1 year



New Single Source Products (Continued) Code 695

For the treatment of patients with clinically significant diabetic macular edema (DME) for whom laser photocoagulation is also indicated; and a hemoglobin A1c of less than 12 percent.

Patients receiving concurrent administration of other anti-VEGF intravitreal injections are not eligible for reimbursement.

Treatment with anti-VEGF agents should only be continued in patients who maintain adequate response to therapy.

Coverage will be provided for patients responding to therapy with another anti-VEGF agent who switch to Eylea HD. Coverage will not be provided for patients who have failed to respond to other anti-VEGF agents.

Recommended Dose: Treatment should be initiated with a monthly intravitreal injection for the first 3 consecutive doses, followed by one injection every 8 to 16 weeks.

LU Authorization Period: 1 year



New Multi-Source Products

Where applicable, please consult the respective brand reference product's drug profile on the ODB e-Formulary for the details of the Limited Use (LU) code and criteria, and/or any associated Therapeutic Notes (TN).

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02548879	Apixaban	2.5mg	Tab	SAI	0.4084
02548887	Apixaban	5mg	Tab	SAI	0.4084

(Interchangeable with Eliquis – GB)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02351102	Famotidine	20mg	Tab	SAI	0.2658
02351110	Famotidine	40mg	Tab	SAI	0.4834

(Interchangeable with Pepcid – GB)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02429772	Jamp Gliclazide-MR	60mg	ER Tab	JPC	0.0632

(Interchangeable with Diamicron MR – GB)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02545772	Mar-Ethosuximide	250mg	Cap	MAR	0.2628

(Interchangeable with Zarontin – GB)



New Multi-Source Products (Continued)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02524090	Mint-Ondansetron Solution	4mg/5mL	Oral Sol	MIN	0.7952

(Interchangeable with Zofran – LU)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02547171	Odan-Ethosuximide	250mg	Сар	ODN	0.2628

(Interchangeable with Zarontin – GB)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02549190	Odan-Valproic Acid	250mg/5mL	Sol	ODN	0.0398

(Interchangeable with Depakene – GB)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02546949	Mint-Varenicline	0.5mg	Tab	MIN	0.4618
02546957	Mint-Varenicline	1 mg	Tab	MIN	0.4618

(Interchangeable with Champix – LU)



New Off-Formulary Interchangeable (OFI) Products

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02549883	Apo-Bilastine Tablets	20mg	Tab	APX	1.1240
02544369	GLN-Bilastine	20mg	Tab	GLP	1.1243
02546795	Jamp Bilastine	20mg	Tab	JPC	1.0217
02547279	Mar-Bilastine	20mg	Tab	MAR	1.1240
02548410	M-Bilastine	20mg	Tab	MAT	1.1240
02551934	NRA-Bilastine	20mg	Tab	NRA	1.1243
02536269	Sandoz Bilastine	20mg	Tab	SDZ	1.1240

(Interchangeable with Blexten)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02551810	Apo-Vilazodone	10mg	Tab	APX	3.0482
02551829	Apo-Vilazodone	20mg	Tab	APX	3.0482
02551837	Apo-Vilazodone	40mg	Tab	APX	4.0577

(Interchangeable with Viibryd)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02430517	Citalopram	10mg	Tab	JPC	0.4464
02409003	Natco-Citalopram	10mg	Tab	NAT	0.4464

(Interchangeable with Celexa)



New Off-Formulary Interchangeable (OFI) Products (Continued)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02547562	Icatibant Injection	30mg/3mL	Inj Sol-Pref Syr	JPC	2025.0000
			3mL Pk		

(Interchangeable with Firazyr)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02545357	Jamp Sumatriptan	50mg	Tab	JPC	9.0650
02545365	Jamp Sumatriptan	100mg	Tab	JPC	9.9866

(Interchangeable with Imitrex)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02531887	Mint-Memantine	10mg	Tab	MIN	1.6357

(Interchangeable with Ebixa)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02550377	NRA-Prucalopride	1mg	Tab	NRA	2.0145
02550385	NRA-Prucalopride	2mg	Tab	NRA	3.1025

(Interchangeable with Resotran)



New Off-Formulary Interchangeable (OFI)
Products (Continued)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02512777	Sandoz AFAtinib	20mg	Tab	SDZ	58.1088
02512785	Sandoz AFAtinib	30mg	Tab	SDZ	58.1088
02512793	Sandoz AFAtinib	40mg	Tab	SDZ	58.1088

(Interchangeable with Giotrif)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	Unit Cost
02330474	Sandoz Amlodipine	2.5mg	Tab	SDZ	0.0767

(Interchangeable with Norvasc)



New Nutrition Products

G.2 PEDIATRIC FORMULA, CHEMICALLY DEFINED – MONOMERIC (ELEMENTAL) MAXIMUM=35.15

Product Name	Strength, Dosage Form, Package Size	PIN/NPN	Mfr	Cost (\$) Per 1000 Kcal	Cost (\$) Per Pkg	Amt (\$) MOH Pays	Amt (\$) Patient Pays
Essential Care Junior (Unflavored)	1.0 kcal/mL Pd- 400g Pouch Pk	09858354	CAM	29.24	57.78	57.78	0.00

H. PEDIATRIC FORMULA, OTHERS

MAXIMUM = N/A

Product Name	Strength, Dosage Form, Package Size	PIN/NPN	Mfr	Cost (\$) Per 1000 Kcal	Cost (\$) Per Pkg	Amt (\$) MOH Pays	Amt (\$) Patient Pays
Renastep	2 kcal/mL-200mL Pk bottle	09858353	VIT	23.08	9.23	9.23	0.00



Transition from Off-Formulary Interchangeable (OFI) to Limited Use (LU)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP
02245619	Copaxone	20mg/mL	Inj Sol Pref Syr- 1mL Pk	TEI	50.6522
02541440	Glatiramer Acetate Injection	20mg/mL	Inj Sol Pref Syr- 1mL Pk	MYL	27.8587

Reason For Use Code and Clinical Criteria

Code 691

As monotherapy for the treatment of patients with relapsing remitting multiple sclerosis (RRMS) meeting ALL the following criteria:

- Recent neurological examination consistent with the diagnosis of RRMS; AND
- Lesions typical of multiple sclerosis on brain magnetic resonance imaging (MRI);
 AND
- Experienced at least 2 clinical attacks in their lifetime with one attack occurring within the prior year; AND
- EDSS score less than or equal to 6.0 prior to start of treatment; AND
- Prescribed by a neurologist who is experienced in the treatment of Multiple Sclerosis.

Note: Transition from another Disease Modifying therapy (DMT) is permitted in those who are deemed to have met the above criteria prior to initiation of the other DMT and if glatiramer acetate is used as monotherapy.

LU Authorization Period: 1 year

Transition from Off-Formulary Interchangeable (OFI) to Limited Use (LU) (Continued)



Code 692

As monotherapy for the treatment of patients who have experienced a single demyelinating event/ Clinically Isolated Syndrome (CIS) meeting ALL the following criteria:

- CIS occurred within the prior 12 months; AND
- Recent neurological examination; AND
- Lesions typical of CIS confirmed on brain magnetic resonance imaging (MRI);
 AND
- EDSS score less than or equal to 6.0 prior to start of treatment; AND
- Prescribed by a neurologist who is experienced in the treatment of Multiple Sclerosis

Note: Transition from another Disease Modifying therapy (DMT) is permitted in those who are deemed to have met the above criteria prior to initiation of the other DMT and if glatiramer acetate is used as monotherapy.

LU Authorization Period: 1 year

Code 693

Renewal of therapy for patients diagnosed with relapsing remitting multiple sclerosis (RRMS) or a single demyelinating event /Clinically Isolated Syndrome (CIS) who meet ALL the following criteria:

- Used as monotherapy for the treatment of RRMS or CIS; AND
- EDSS score less than or equal to 6.0; AND
- Disease activity is stabilized as determined by a neurological exam and the number of clinical relapses experienced while on treatment; AND
- Prescribed by a neurologist experienced in the treatment of Multiple Sclerosis (MS) OR a prescriber in consultation with a neurologist overseeing the patient's MS.

LU Authorization Period: 1 year

Limited Use Code & Clinical Criteria Changes

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02245397	NovoRapid	100U/mL	Inj Sol-10mL Pk	NOO

Codes 642, 643: LU codes and criteria ended as of the December 2024 formulary update.

Code 646: This LU code and criteria remain unchanged.

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02244353	NovoRapid Penfill*	100U/mL	Inj Sol-5x3mL Pk	NOO
02377209	NovoRapid FlexTouch*	100U/mL	Inj Sol-Prefil 5X3mL Pk	NOO
			Disposable Pen	

Codes 642, 643: LU code and criteria ended as of the December 2024 formulary update.

*Products delisted from the formulary effective December 2024 formulary update.

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02245689	Lantus-(Vial)	100U/mL	Inj Sol-10mL Vial Pk	SAV

Codes 642, 643: LU codes and criteria ended as of the December 2024 formulary update.

Code 644: This LU code and criteria remain unchanged.



Limited Use Code & Clinical Criteria Changes (Continued)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02294338	Lantus Solostar*	100U/mL	Inj Sol-5x3mL Pk	SAV
02251930	Lantus-(Cartridge)*	100U/mL	Inj Sol-5x3mL Pk	SAV

Codes 642, 643: LU code and criteria ended as of the December 2024 formulary update.

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02229704	Humalog	100U/mL	Inj Sol-10mL Pk	LIL

Codes 642, 643: LU code and criteria ended as of the December 2024 formulary update.

Code 646: This LU code and criteria remain unchanged.

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02403412	Humalog Kwikpen*	100U/mL	Inj Sol-5x3mL Pk	LIL
09853715	Humalog*	100U/mL	Inj Sol-5x3mL Pk	LIL
02470152	Humalog*	100U/mL	Inj Sol-Pref Pen 5x3mL Pk	LIL
			(Junior KwikPen)	

Codes 642, 643: LU code and criteria ended as of the December 2024 formulary update.

^{*}Products delisted from the formulary effective December 2024 formulary update.

^{*}Products delisted from the formulary effective December 2024 formulary update.



Revision of Limited Use Criteria

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02525267	Bimzelx	160mg/mL	Inj Sol-1mL Pref Syr	UCB
00505075	D: 1	400 / 1	(Preservative-Free)	1100
02525275	Bimzelx	160mg/mL	Inj Sol-1mL Pref Autoinj (Preservative-Free)	UCB

Revised Clinical Criteria

Code 641

The existing paragraph on approvals is replaced by the paragraph below. No other changes to the criteria.

Approvals will allow for standard dosing for Bimzelx, which is 320mg subcutaneously every 4 weeks for the first 16 weeks, and every 8 weeks thereafter. A dose of 320mg every 4 weeks after the first 16 weeks may be considered in patients with a body weight of 120kg or more who did not achieve a complete skin response. For patients with no improvement after 16 weeks of treatment at the Health Canada approved dose, higher or more frequent doses are not recommended and the physician should consider switching to an alternative biologic agent.

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02480522	Xydalba	500mg/Vial	Pd for Sol (Preservative-Free)	EVL

Code 677

The existing paragraph is replaced by the paragraph below. No other changes to the criteria.

4. Patient is deemed to be at high risk of nonadherence to a standard antibiotic treatment for MRSA ABSSSI or treatment with dalbavancin rather than a standard antibiotic treatment for MRSA ABSSSI will avoid the need for hospitalization OR will limit the need for prolonged hospitalization; AND

Revision of Limited Use Criteria (Continued) Ontario



DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02343541	Prolia (Preservative Free)	60mg/mL	Inj Sol-Pref Syr	AMG

Code 690

Only for a patient who is established on Prolia (denosumab) therapy prior to November 29, 2024 and who is or becomes palliative requiring end-of-life care during the biosimilar transition period between November 29, 2024 to August 29, 2025.

Patient must also meet the following criteria:

- The patient is a postmenopausal female or male with osteoporosis and Prolia (denosumab) is being used to increase the bone mass in the patient;
- The patient is at high risk* for fracture; and
- One of the following applies to the patient:
 - o The patient has failed other available osteoporosis therapy (i.e., fragility fracture or evidence of a decline in bone mineral density below pretreatment baseline levels) despite adherence to the therapy for one year; or
 - bisphosphonates are contraindicated for the patient due to hypersensitivity, abnormalities of the esophagus (e.g., esophageal stricture or achalasia), or inability to stand or sit upright for at least 30 minutes.

*High risk of fracture is defined as one of the following:

- o a prior fragility fracture AND a moderate 10-year fracture risk (10% to 20%) based on the Canadian Association of Radiologists and Osteoporosis Canada (CAROC) tool or the Fracture Risk Assessment (FRAX) tool;
- o a high 10-year fracture risk (greater than or equal to 20%) based on the CAROC or FRAX tool: or
- where a patient's 10-year fracture risk based on the CAROC or FRAX tool is less than the thresholds defined above, a high fracture risk based on evaluation of clinical risk factors for fracture.

Note: Use of the CAROC or FRAX tool may underestimate fracture risk in certain circumstances and may not include all risk factors.

Funded duration: One period of up to 12 months beginning on the date of the first prescription with RFU code 690 is dispensed for the patient.

NOTE: In all cases, patients receiving Prolia must not be receiving concomitant bisphosphonate therapy. The recommended dose of PROLIA (denosumab) is a single SC injection of 60 mg, once every 6 months.

LU Authorization Period: 12 months from date of authorization.



Manufacturer Name Changes

DIN/PIN	Product Name	Strength	Dosage Form	Current Mfr	New Mfr
02459418	Revlimid	2.5mg	Сар	CEL	BQU
02304899	Revlimid	5mg	Сар	CEL	BQU
02304902	Revlimid	10mg	Сар	CEL	BQU
02317699	Revlimid	15mg	Сар	CEL	BQU
02440601	Revlimid	20mg	Сар	CEL	BQU
02317710	Revlimid	25mg	Сар	CEL	BQU



Product Name and Manufacturer Name Changes

DIN/PIN	Current Product Name	Current Mfr	New Product Name	New Mfr	Strength	Dosage Form
02255529	Co Fluvoxamine	COB	Teva-Fluvoxamine	TEV	50mg	Tab
02255537	Co Fluvoxamine	COB	Teva-Fluvoxamine	TEV	100mg	Tab



Drug Benefit Price (DBP) Changes

DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP/Unit
					Price
00771376	AA-Diltiaz	30mg	Tab	AAP	0.2168
02231329	Apo-Fluvoxamine	50mg	Tab	APX	0.5410
02231330	Apo-Fluvoxamine	100mg	Tab	APX	0.9728
02291967	Apo-Ondansetron	4mg/5mL	O/L	APX	0.7952
02243324	Apo-Propafenone	150mg	Tab	APX	0.6306
02255529	Co Fluvoxamine	50mg	Tab	COB	0.5410
02255537	Co Fluvoxamine	100mg	Tab	COB	0.9728
02244126	Dovobet	50mcg/g &	Oint	LEO	2.1425
		0.5mg/g			
02319012	Dovobet Gel	50mcg/g &	Top Gel	LEO	2.1375
		0.5mg/g	_	_	
01976133	Dovonex	50mcg/g	Oint	LEO	1.2208
02270811	Finacea	15%	Top Gel	LEO	0.8566
00586668	Fucidin	2%	Cr	LEO	1.0950
00586676	Fucidin	2%	Oint	LEO	1.0950
09857367	Innohep	2500IU/0.25mL	Inj Pref Syr	LEO	7.1400
02358158	Innohep	3500IU/0.35mL	Inj Pref Syr	LEO	9.9860
02358166	Innohep	4500IU/0.45mL	Inj Pref Syr	LEO	12.8420
02429462	Innohep	8000IU/0.4mL	Inj Pref Syr	LEO	22.2100
02231478	Innohep	10000IU/0.5mL	Inj Pref Syr	LEO	29.1240
02429470	Innohep	12000IU/0.6mL	Inj Pref Syr	LEO	34.9820
02358174	Innohep	14000IU/0.7mL	Inj Pref Syr	LEO	40.8110
02429489	Innohep	16000IU/0.8mL	Inj Pref Syr	LEO	46.6430
02358182	Innohep	18000IU/0.9mL	Inj Pref Syr	LEO	52.4660
02167840	Innohep	10000IU/mL	Inj-2mL Pk	LEO	53.9070
02229515	Innohep	20000IU/mL	Inj-2mL Pk	LEO	114.9800
02539411	Jamp Amoxi Clav	200mg &	Susp	JPC	0.1007
	Suspension	28.5mg/5mL			
02542587	Jamp Ipratropium HFA	20mcg/Metered	Inh-200 Dose	JPC	11.2530
		Dose	Pk		
02490617	Jamp Ondansetron	4mg/5mL	O/L	JPC	0.7952
02517469	Jamp Quinapril	20mg	Tab	JPC	0.4642
02517477	Jamp Quinapril	40mg	Tab	JPC	0.4642

Drug Benefit Price (DBP) Changes (Continued)



DIN/PIN	Product Name	Strength	Dosage Form	Mfr	DBP/Unit Price
02549433	Mar-Eszopiclone	1mg	Tab	MAR	1.4365
02549441	Mar-Eszopiclone	2mg	Tab	MAR	1.4365
02549468	Mar-Eszopiclone	3mg	Tab	MAR	1.4365
02457172	Mylan-Propafenone	150mg	Tab	MYL	0.6306
09853588	Peptamen Junior		Liq-250mL Pk	NES	8.8150
09857562	Peptamen Junior 1.5	1.5Kcal/mL	Liq - 250mL Tetra Pk	NES	13.2224
09857101	Peptamen with Prebio	1kcal/mL	Liq-250mL Pk	NES	8.8150
09857102	Peptamen with Prebio	1kcal/mL	Liq-1500mL Pk	NES	52.8900
02324253	PMS-Hydrocodone	1mg/mL	O/L	PMS	0.0617/mL
02340577	PMS-Quinapril	20mg	Tab	PMS	0.4642
02340585	PMS-Quinapril	40mg	Tab	PMS	0.4642
09854401	Portagen	1.02kcal/mL	Pd-454g Pk	MJN	55.7800
02244148	Protopic	0.1%	Oint	LEO	3.5866
02244149	Protopic	0.03%	Oint	LEO	3.3530
09857369	PurAmino A+	5kcal/g	Pd-400g Pk	MJN	55.7900
02371081	Xeomin	50 LD50 Units	Pd for Inj-Vial Pk	MEZ	169.4550
02324032	Xeomin	100 LD50 Units	Pd for Inj-Vial Pk	MEZ	338.9100



Discontinued Product

(Some products will remain on Formulary for six months to facilitate depletion of supply)

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02132664	Fragmin	10000IU/mL	Inj Sol-1mL Pk	PFI
02262800	Strattera	10mg	Сар	LIL
02262819	Strattera	18mg	Сар	LIL
02262827	Strattera	25mg	Сар	LIL
02262835	Strattera	40mg	Сар	LIL
02262843	Strattera	60mg	Сар	LIL



Delisted Products

DIN/PIN	Product Name	Strength	Dosage Form	Mfr
02464276	Adlyxine	0.05mg/mL	Inj Sol-Pref Pen 3mL Pk	SAC
02464284	Adlyxine	0.1mg/mL	Inj Sol-Pref Pen 3mL Pk	SAC
02248501	Apo-Quinapril	20mg	Tab	APX
02248502	Apo-Quinapril	40mg	Tab	APX
02221861	Anandron	50mg	Tab	CHE
02291177	Champix	0.5mg	Tab	PFI
02291185	Champix	1.0mg	Tab	PFI
09857519	Champix	0.5mg &	Tab (Starter Pack-53	PFI
		1.0mg	Tabs)	
02287153	Fosrenol	500mg	Chew Tab	TAK
02287161	Fosrenol	750mg	Chew Tab	TAK
02287188	Fosrenol	1000mg	Chew Tab	TAK
09857633	FreeStyle Libre Reader			ABD
	Flash Glucose Monitoring			
	System - Reader			
09857632	FreeStyle Libre Sensor			ABD
	Flash Glucose Monitoring			
	System - Sensor			
09853715	Humalog	100U/mL	Inj Sol-5x3mL Pk	LIL
02403412	Humalog Kwikpen	100U/mL	Inj Sol-5x3mL Pk	LIL
02470152	Humalog	100U/mL	Inj Sol-Pref Pen 5x3mL	LIL
00004000		40011/	Pk (Junior KwikPen)	0.41.7
02294338	Lantus Solostar	100U/mL	Inj Sol-5x3mL Pk	SAV
02251930	Lantus-(Cartridge)	100U/mL	Inj Sol-5x3mL Pk	SAV
02489368	Latanoprost&Timolol	50mcg/mL &	Oph Sol-2.5mL Pk	TCI
00044252	Ophthalmic Solution	5mg/mL	Ini Cal Ev2ml Dk	NOO
02244353	NovoRapid Penfill	100U/mL	Inj Sol-5x3mL Pk	NOO
02377209	NovoRapid FlexTouch	100U/mL	Inj Sol-Prefil 5X3mL Pk Disposable Pen	NOO
02294559	PMS-Propafenone	150mg	Tab	PMS
02343053	Propafenone	150mg	Tab	SAI
00862924	Teva-Diltazem	30mg	Tab	TEV
09853308	Vivonex Pediatric		Pd-48.7g Pk	NES

